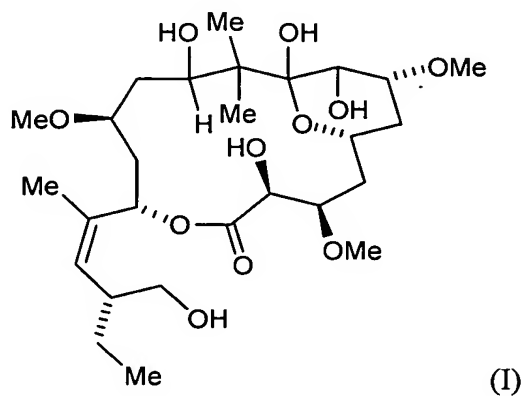


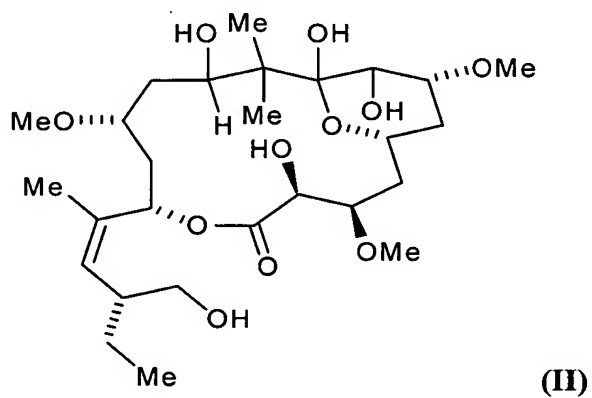
AMENDMENTS TO THE CLAIMS

This listing replaces all prior versions and listings of claims in the application.

1. (Original) A synthetic compound of formula:



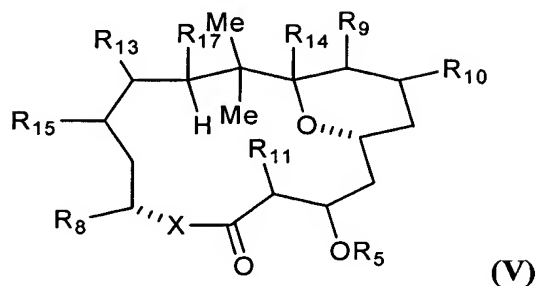
2. (Original) A synthetic compound of formula:



(III)

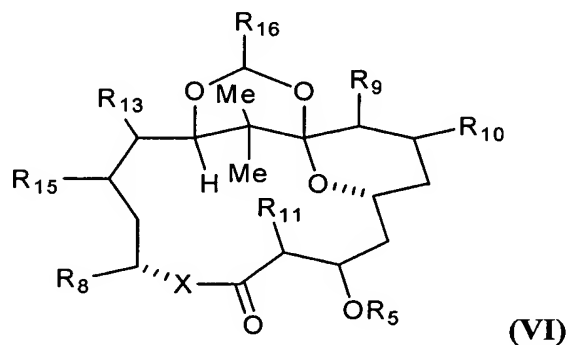
4. (Original) A compound of formula:

5. (Original) A compound of formula:



where R_{13} is H or Me, where R_{14} , R_{17} can be the same or different and are selected from the group consisting of H, OH, and OR, where R_9 , R_{10} , R_{11} , R_{15} can be the same or different and are selected from the group consisting of H, Me, and OR, where R and R_5 can be the same or different and are selected from the group consisting of H, Me, alkyl, or functionalized alkyl, where R_8 is selected from the group consisting of H, aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, where X is O or NH, and wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

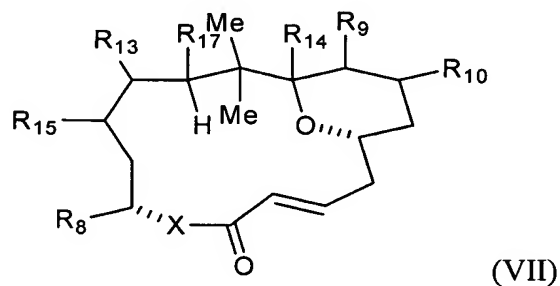
6. (Original) A composition comprising a compound of formula:



where R_{13} is H or Me, where R_9 , R_{10} , R_{11} , R_{15} can be the same or different and are selected from the group consisting of H, Me, OR, where R and R_5 can be the same or different and are selected from the group consisting of H, Me, alkyl, or functionalized alkyl, where R_8 , R_{16} can be the same or different and are selected from the group consisting of H, aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized

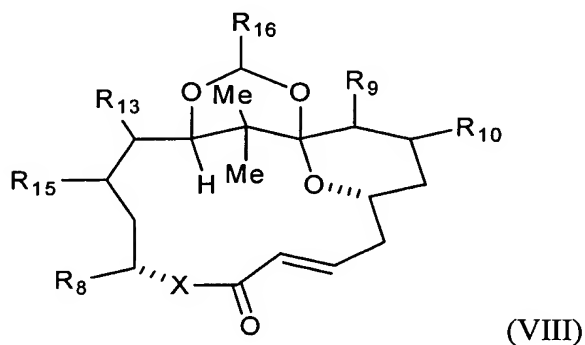
alkynyl, and where X is O or NH and wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

7. (Original) A composition comprising a compound of formula:



where R₁₃ is H or Me, where R₁₄, R₁₇ can be the same or different and is selected from the group consisting of H, OH, or OR, where R₉, R₁₀, R₁₅ can be the same or different and are selected from the group consisting of H, Me, OR, where R is selected from the group consisting of H, Me, alkyl, or functionalized alkyl, where R₈ is selected from the group consisting of H, aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, where X is O or NH, and wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

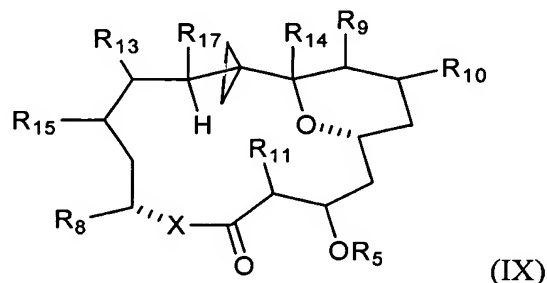
8. (Original) A composition comprising a compound of formula:



where R₁₃ is H or Me, where R₉, R₁₀, R₁₅ can be the same or different and are selected from the group consisting of H, Me, OR, where R is selected from the group consisting of H, Me, alkyl, or functionalized alkyl, where R₈, R₁₆ can be the same or different and are selected

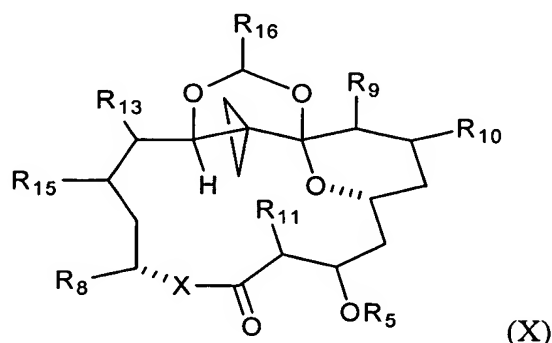
from the group consisting of H, aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, where X is O or NH, and wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

9. (Original) A composition comprising a compound of formula:



where R₁₃ is H or Me, where R₁₄, R₁₇ can be the same or different and are selected from the group consisting of H, OH, or OR, where R₉, R₁₀, R₁₁, R₁₅ can be the same or different and are selected from the group consisting of H, Me, OR, where R and R₅ can be the same or different and are selected from the group consisting of H, Me, alkyl, or functionalized alkyl, where R₈ is H, aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, where X is O or NH, and wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

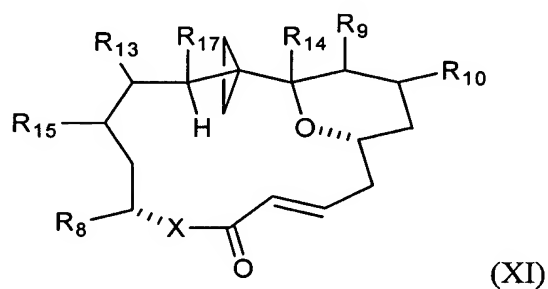
10. (Original) A composition comprising a compound of formula:



where R₁₃ is H or Me, where R₉, R₁₀, R₁₁, R₁₅ can be the same or different and are selected from the group consisting of H, Me, OR, where R and R₅ can be the same or different and are

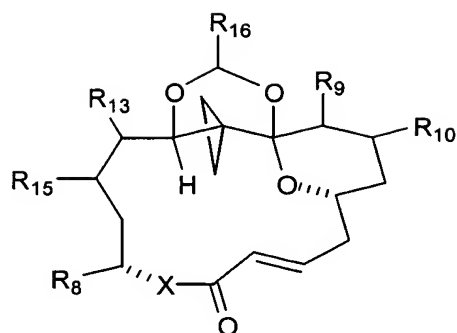
selected from the group consisting of H, Me, alkyl, or functionalized alkyl, where R_8 , R_{16} can be the same or different and are selected from the group consisting of H, aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, where X is O or NH, and wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

11. (Original) A composition comprising a compound of formula:



where R_{13} is H or Me, where R_{14} , R_{17} can be the same or different and are selected from the group consisting of H, OH, or OR, where R_9 , R_{10} , R_{15} can be the same or different and are selected from the group consisting of H, Me, OR, where R is selected from the group consisting of H, Me, alkyl, or functionalized alkyl, where R_8 is H, aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, where X is O or NH, and wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

12. (Original) A composition comprising a compound of formula:



(XII)

where R₁₃ is H or Me, where R₉, R₁₀, R₁₅ can be the same or different and are selected from the group consisting of H, Me, OR, where R is selected from the group consisting of H, Me, alkyl, or functionalized alkyl, where R₈, R₁₆ can be the same or different and are selected from the group consisting of H, aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, where X is O or NH, and wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

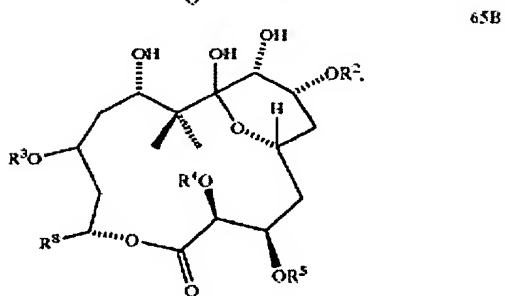
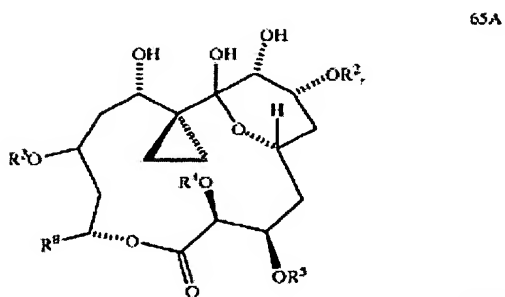
13. (Original) A synthetic compound having the ¹³C NMR signature of **FIG. 4** and the ¹H NMR signature of **FIG. 5**, wherein the compound is dextrarotary, and wherein the compound comprises microtubule-stabilizing activity.
14. (Original) A method for treating cancer comprising contacting a tumor cell within a subject with a compound of any one of claims 1 through 13 for a period of time and in an amount sufficient to inhibit growth of the tumor cell.
15. (Original) A method of suppressing growth of a tumor cell comprising contacting said cell with a compound of any one of claims 1 through 13 for a period of time and in an amount sufficient to suppress growth of the tumor cell.
16. (Original) A method of inhibiting growth of proliferating cells comprising the step of administering to the proliferating cells the compound of any one of claims 1

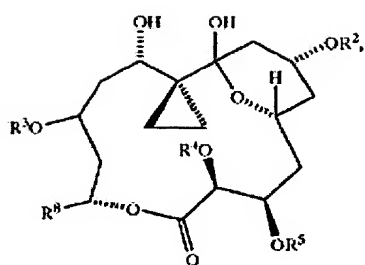
through 13 for a period of time and in an amount sufficient to inhibit proliferation of the cells.

17. (Original) A method of stabilizing microtubule formation in a cell comprising administering to the cell the compound of any one of claims 1 through 13 for a period of time and in an amount sufficient to stabilize microtubule formation.

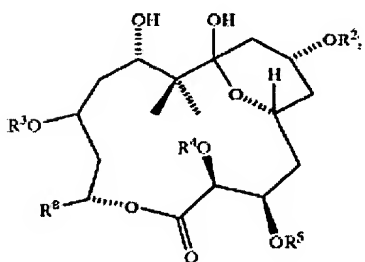
18-22. (Canceled)

23. (Original) A compound selected from the group consisting of:

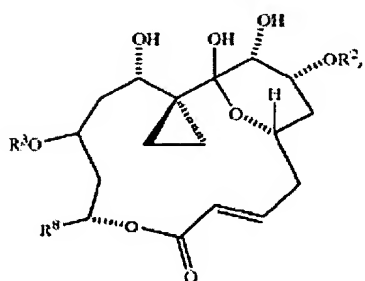




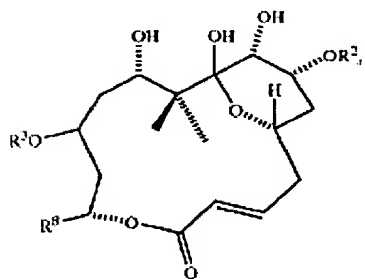
65C



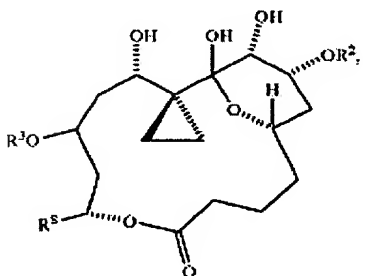
65D



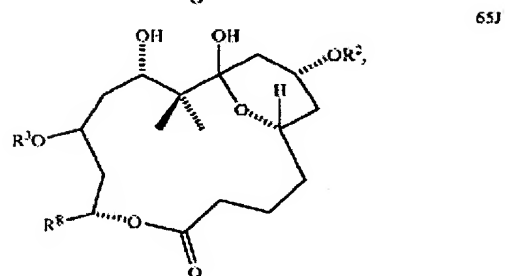
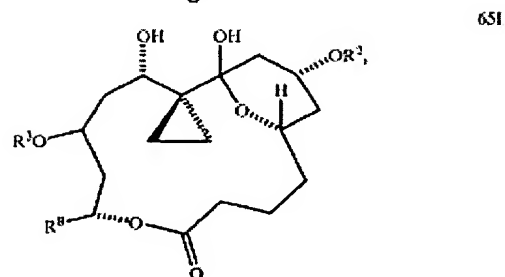
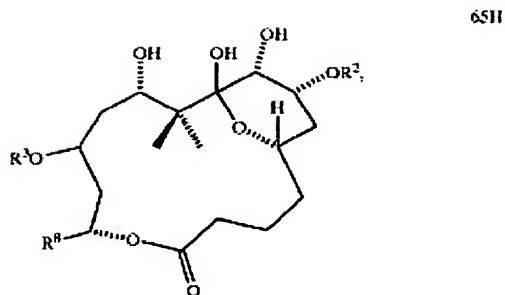
65E



65F

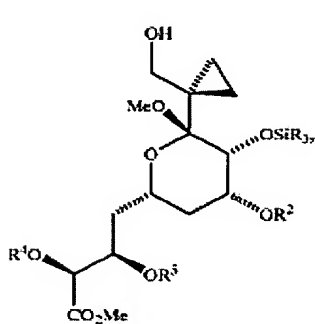


65G



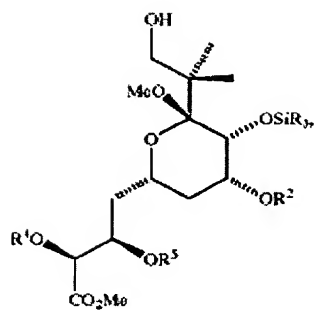
wherein R^2 , R^3 , R^4 , R^5 can be the same or different and are selected from the group consisting of alkyl, and functionalized alkyl, and where R^8 is selected from the group consisting of aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

24. (Original) A method of producing Peloruside A comprising:
- (a) oxidizing an alcohol function in a compounds selected from the group consisting of:

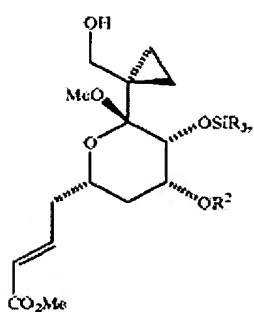


61A

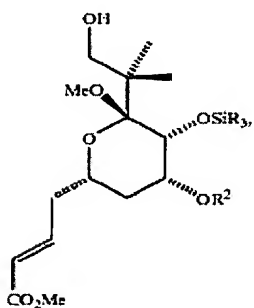
61B



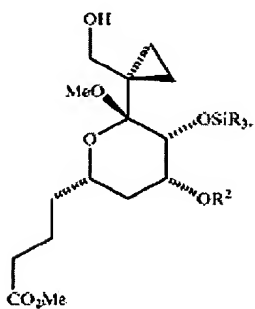
61E



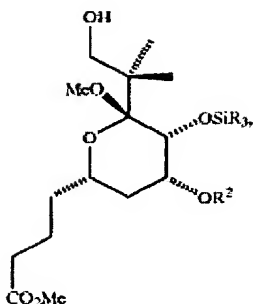
61F



61G



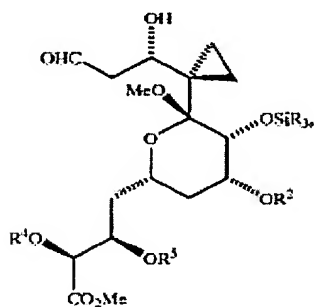
61H



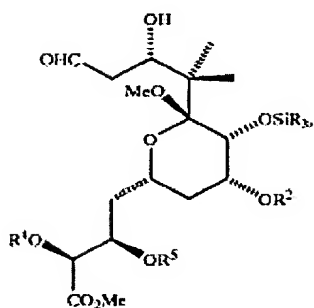
wherein R^2 , R^4 and R^5 are the same or different and are selected from the group consisting of alkyl, and functionalized alkyl, wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl, to obtain a compound having an aldehyde function;

- (b) reacting the compound obtained in (a) with an allylating agent;
- (c) subjecting the reaction product of (b) to oxidative cleavage of the aldehyde to obtain a compound selected from the group consisting of:

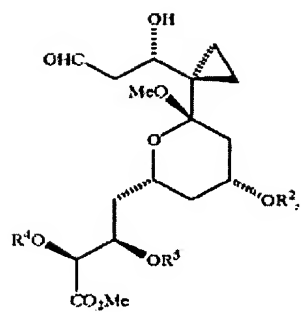
62A



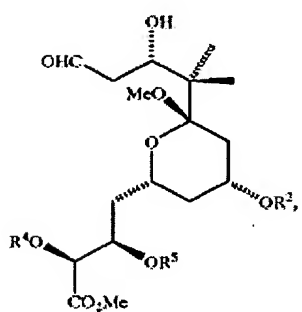
62B



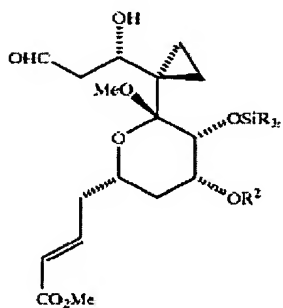
62C



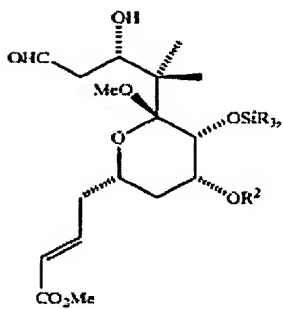
62D

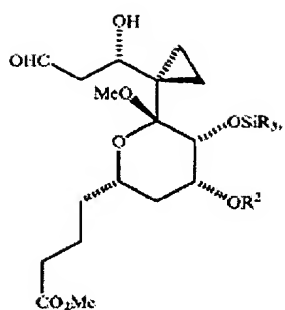


62E

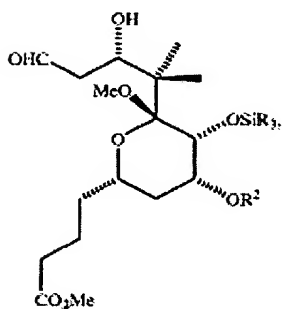


62P

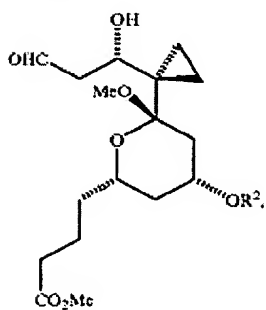




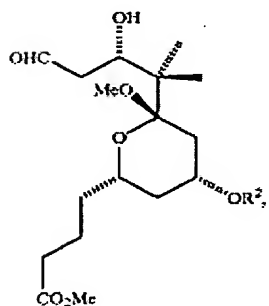
62G



62H



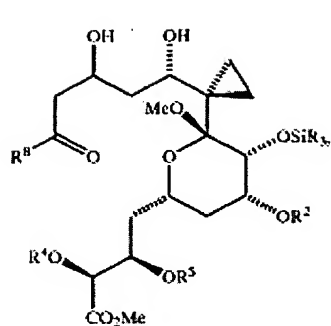
62I



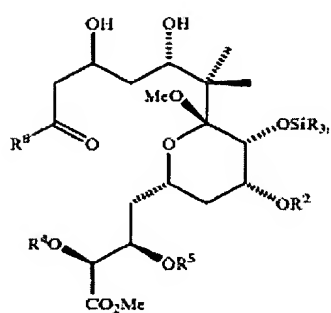
62J

wherein R^2 , R^4 and R^5 are the same or different and are selected from the group consisting of alkyl, and functionalized alkyl, wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl;

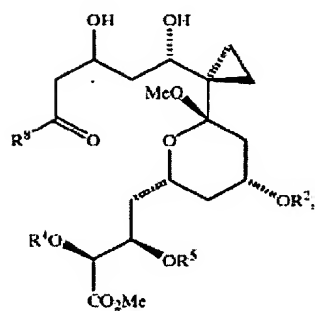
(d) reacting the compound obtained in (c) with an enolate derived from a methyl or ethyl ketone to obtain a compound selected from the group consisting of:



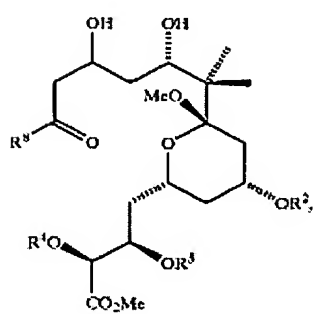
63A



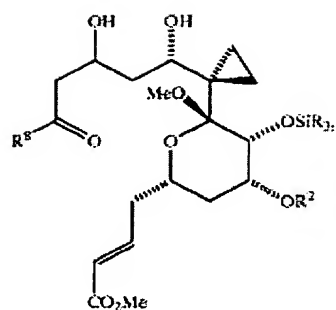
63B



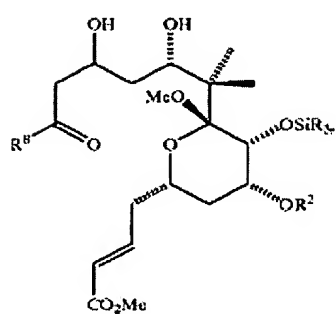
63C



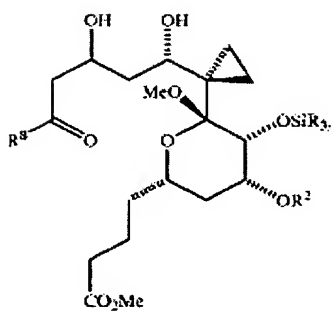
63D



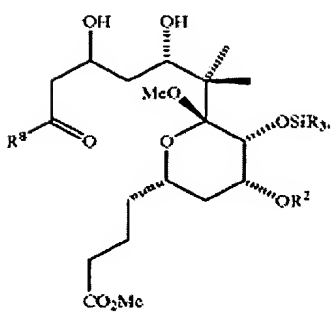
63E



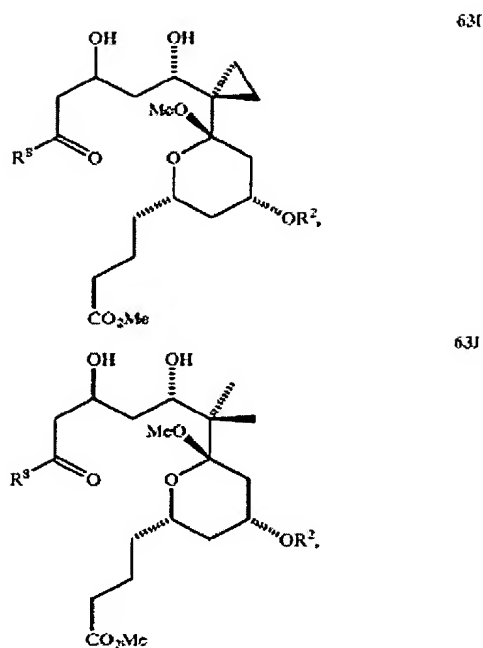
63F



63G

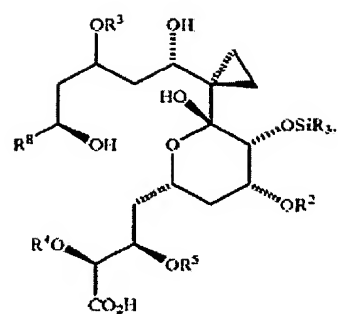


63H

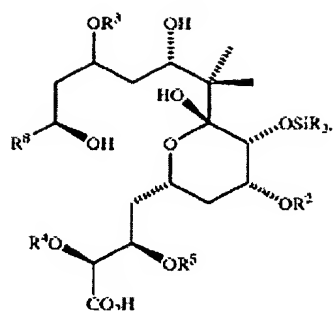


wherein R^2 , R^4 and R^5 can be the same or different and are selected from the group consisting of alkyl, and functionalized alkyl, and R^8 is selected from the group consisting of aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl;

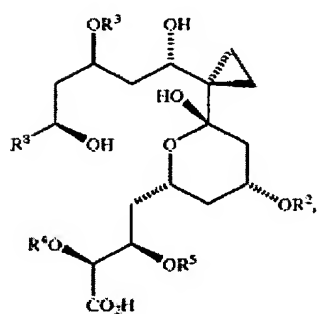
- (e) reacting the compound obtained in (d) with an alkylating agent to introduce and R^3 group;
- (f) subjecting the compound obtained in (e) with a reducing agent to reduce the ketone group in the compound obtained in (e) to an alcohol;
- (g) converting the alcohol group of the compound obtained in (f) to an ester group;
- (h) subjecting the compound obtained in (g) to an agent that hydrolyzes the ester group of the compound produced in (f) to a carboxylic acid group to obtain a seco-acid compound selected from the group consisting of:



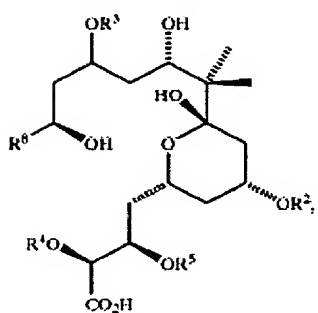
64A



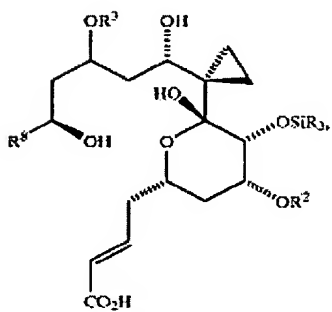
64B



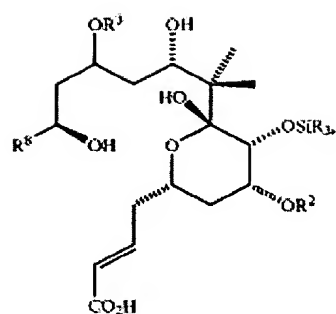
64C



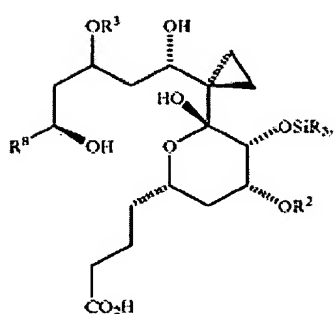
64D



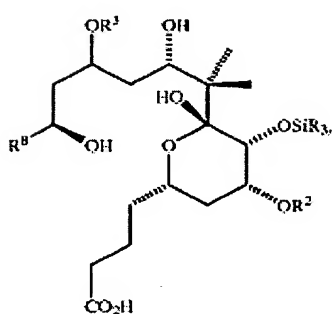
64E



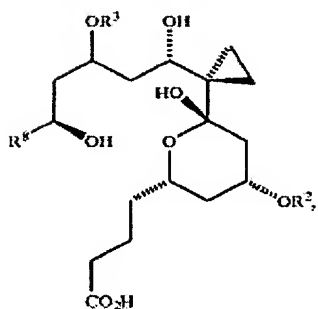
64F



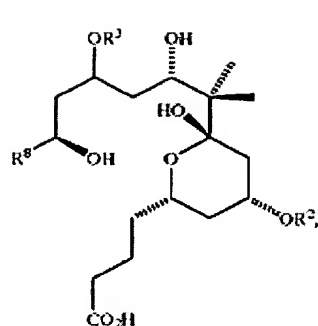
64G



64H

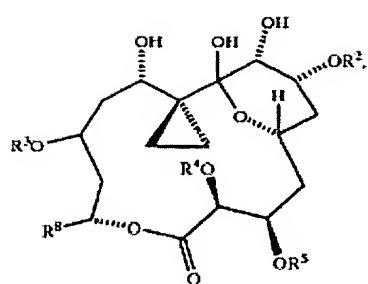


64I

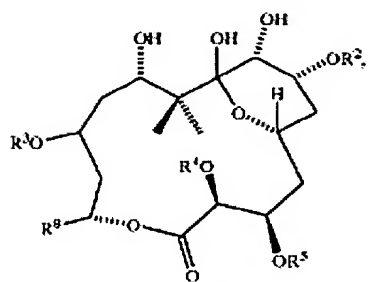


wherein R², R³, R⁴, R⁵ can be the same or different and are selected from the group consisting of alkyl, and functionalized alkyl, and where R⁸ is selected from the group consisting of aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl; and

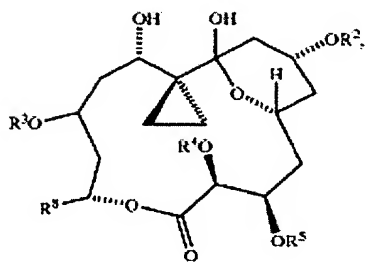
- (i) reacting the carboxylic acid group and a hydroxyl group of the compound produced in (h) to obtain a macrolactone selected from the group consisting of:



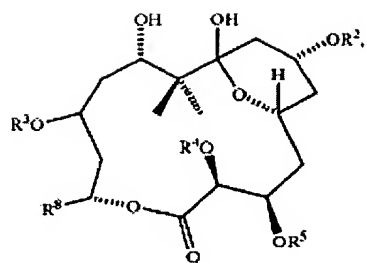
65A



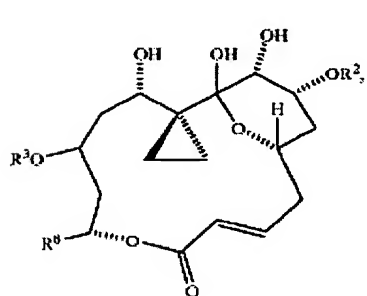
65B



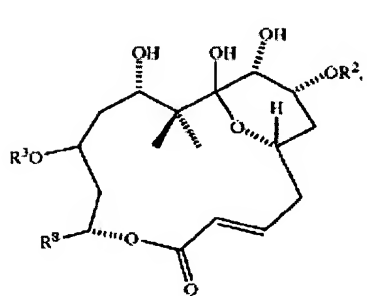
65C



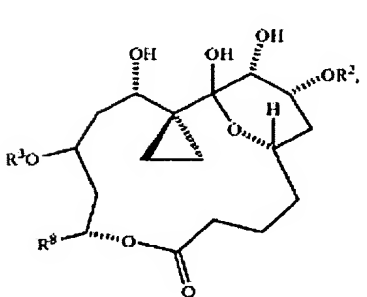
65D



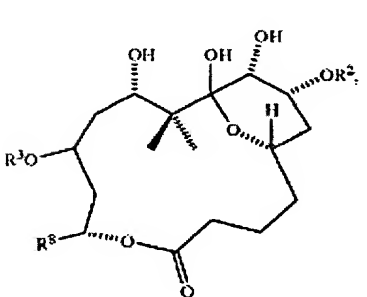
65E



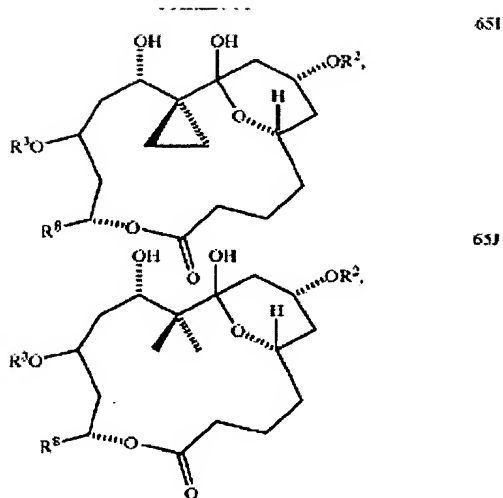
65F



65G

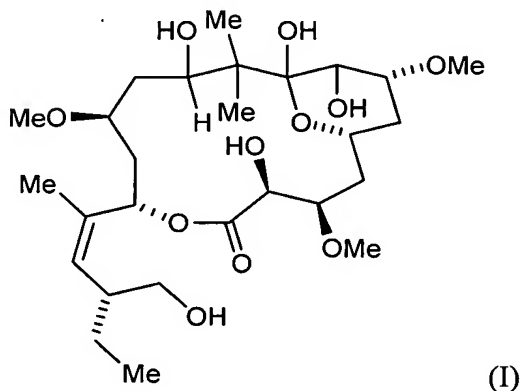


65H



wherein R^2 , R^3 , R^4 , R^5 can be the same or different and are selected from the group consisting of alkyl, and functionalized alkyl, and where R^8 is selected from the group consisting of aryl, heteroaryl, alkyl, functionalized alkyl, alkenyl, functionalized alkenyl, alkynyl, and functionalized alkynyl, wherein the functional group is a heteroatom, a halide, an aryl, or a heteroaryl.

25. (Original) The method of claim 24 wherein the macrolactone is Peloruside A.
26. (Original) The method of claim 24 wherein the macrolactone has the formula:



27. (Original) The method of claim 24 wherein the macrolactone has the formula:

